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		Elizabeth Peet Sabino / Hollis N Erb / James L Catalfamo, J Am Vet Med Assoc, Feb 2006	We did this by add quotes to common
		activity (CBA) to screen for <b>type II von Willebrand disease</b> (vWD) in dogs. Sample Population-293samples from dogs with inherited <b>type II</b> or III vWD. Procedure-Bovinediscriminate between type I and <b>type II</b> vWD.	and by removing n essential words Repeat without
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$\overline{}$	2	Development of a collagen-binding activity assay as a screening test for	adolescent
LJ	۷.	type II von Willebrand disease in dogs.	base sequence
		Elizabeth Peet Sabino / Hollis N Erb / James L Catalfamo, Am J Vet	blood platelets
		Res, Feb 2006	cofactor activity
		Objective-To develop an assay to measure canine von Willebrand factor (vWF):collagen-binding activity (CBA) to screen for type 2 <b>von Willebrand</b>	<u>collagen</u>
		<b>disease</b> (vWD) in dogs. Sample Population-293 plasma samples submitted	factor antigen
		for analysis of canine vWF antigen (vWF:Ag) and 12 control	<u>haemostasis</u>
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		vious all 00 magnite from MEDI INE (DubMed	molecular sequenc
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	3.	A new variant of von Willebrand disease (type II I) with a normal degree of proteolytic cleavage of von Willebrand factor.  G Castaman / F Rodeghiero / A Lattuada / P M Mannucci, Thromb	polymerase chain i research support, i gov't, p.h.s.
		Res, Feb 1992	von willebrand fact
		A variant of <b>type II von Willebrand disease</b> (vWd) is described in a	multimers
		young woman and her mother with severe lifelong bleeding historiesthis	Or refine using:
		variant of vWd appears to differ from those reported hitherto, the designation of <b>type II</b> I is proposed.	All of the words

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4.	Type II H von Willebrand disease: new structural abnormality of plasma and platelet von Willebrand factor in a patient with  A B Federici / P M Mannucci / R Lombardi / A Lattuada / M L Colibretti / J A Dent / T S Zimmerman, Am J Hematol, Dec 1989 In this study a new variant of type II von Willebrand disease is identified by multimeric analyses of increasing resolving powerdescribed. Therefore the proposed designation for this new variant is type II H.	
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5.	A new variant of type II von Willebrand disease with aberrant multimeric structure of plasma but not platelet von Willebrand  P M Mannucci / R Lombardi / A B Federici / J A Dent / T S  Zimmerman / Z M Ruggeri, Blood, Jul 1986diagnosed as having Type II von Willebrand disease. The larger multimersThis new variant of von Willebrand disease, therefore, is characterizedproposed for this type of von Willebrand disease in accordance with	
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6.	von Willebrand Disease Associated with Superficial Temporal Artery Pseudoaneurysm Ricciardo, B.J. / Mwipatayi, B.P. / Abbas, M. / Sieunarine, K. / Eikelboom, J.W., European Journal of Vascular & Endovascular Surgery, Sep 2005two patients with von Willebrand disease (vWD) that presentedKeywords Pseudoaneurysm von Willebrand disease Collagen Temporal artery von Willebrand disease is a common bleedingIII) or dysfunction (type II) of von Willebrand Published journal article available from	
7.	Multiple normal deliveries in a woman with severe portal hypertension due to congenital hepatic fibrosis: the importance of  Ayse L Mindikoglu / Arie Regev / Mary J O'Sullivan / Eugene R  Schiff, Am J Gastroenterol, Oct 200526-year-old woman with congenital hepatic fibrosis (CHFib), adult polycystic kidney disease, type II diabetes mellitus, and von Willebrand disease, who despite persistent advice to the contrary because of severe portal hypertension, had 5  MEDLINE/PubMed Citation on Publiced	
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8.	A molecular approach to the classification of von Willebrand disease Schneppenheim, R. / Budde, U. / Ruggeri, Z.M., Best Practice &	

Research in Clinical Haematology, Jun 2001 ...B A new variant of type II von Willebrand disease with aberrant multimeric...McKeown L P A variant of type II von Willebrand disease with an abnormal triplet...M A new variant of von Willebrand disease (type II I) with a normal degree... Published journal article available from PCIBNCE @DIRECT view all 61 results from ScienceDirect similar results **9.** Management of inherited von Willebrand disease Mannucci, P.M. / Federici, A.B., Best Practice & Research in Clinical Haematology, Jun 2001 ...vasopressin (DDAVP) in type II B von Willebrand disease New England Journal of...R DDAVP for type IIB von Willebrand disease Blood 74 1989 1859 1860...vasopressin (DDAVP) in type II B von Willebrand disease European Journal of Haematology... Published journal article available from science doinger view all 61 results from ScienceDirect similar results ☐ 10. A molecular approach to the classification of von Willebrand disease Schneppenheim, R. / Budde, U. / Ruggeri, Z.M., Best Practice & Research in Clinical Haematology, Jun 2001 ...B A new variant of type II von Willebrand disease with aberrant multimeric...McKeown L P A variant of type II von Willebrand disease with an abnormal triplet...M A new variant of von Willebrand disease (type II I) with a normal degree... Published journal article available from BCIENCE @DIRECT view all 61 results from ScienceDirect similar results **11.** Management of inherited von Willebrand disease Mannucci, P.M. / Federici, A.B., Best Practice & Research in Clinical Haematology, Jun 2001 ...vasopressin (DDAVP) in type II B von Willebrand disease New England Journal of...R DDAVP for type IIB von Willebrand disease Blood 74 1989 1859 1860...vasopressin (DDAVP) in type II B von Willebrand disease European Journal of Haematology... Published journal article available from SCIENCE @PIRECT view all 61 results from ScienceDirect similar results 12. A new variant of von willebrand disease (type II I) with a normal degree of proteolytic cleavage of von Willebrand... Castaman, G. / Rodeghiero, F. / Lattuada, A. / Mannucci, P.M., Thrombosis Research, Feb 1992 A variant of type II von Willebrand disease (vWd) is described in a young woman and her mother with severe lifelong bleeding histories. On electrophoresis with low-resolution agarose gels the plasma of the proband lacked large and... Published journal article available from SCIENCE @DIRECT view all 61 results from ScienceDirect similar results 13. Identification of a new candidate mutation, G1629R, in a family with type 2A von Willebrand disease. P Casaña / F Martínez / S Haya / J A Aznar, Am J Hematol, Apr 1999 Type 2A is a qualitative variant of **von Willebrand disease** (vWD)

characterized by a reduced platelet-dependent function, associated with an absence of large multimers. A G5135A transition...

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**14.** <u>Influence of mutations and size of multimers in type II von Willebrand</u> disease upon the function of von Willebrand factor.

O Christophe / A S Ribba / D Baruch / B Obert / C Rouault / K Niinomi / G Piétu / (...) / J P Girma, Blood, Jun 1994 ...factor (vWF) from normal individuals and from two patients with type IIA (Glu875Lys) and type IIB (duplication of Met 540) von Willebrand disease (vWD) with the corresponding fully multimerized recombinant proteins. We included cryosupernatant from normal human plasma...

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15. New variant of von Willebrand disease type II with markedly increased levels of von Willebrand factor antigen and dominant...

M R Ledford / I Rabinowitz / J E Sadler / J W Kent / F Civantos, Blood, Jul 1993

A variant of **von Willebrand disease** (vWD) was identified in six members of a kindred spanning four generations. The proband was a 46-year-old woman with a lifelong...

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**16.** A variant of type II von Willebrand disease with an abnormal triplet structure and discordant effects of protease inhibitors...

H R Gralnick / S B Williams / L P McKeown / P Maisonneuve / C Jenneau / Y Sultan, Am J Hematol, Mar 1987

...characterized the plasma and platelet von Willebrand factor (vWf) multimeric structure of a patient with **von Willebrand disease** (vWd) as having a long bleeding time, no aggregation of her platelet-rich plasma (PRP) to ristocetin, and...

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17. von Willebrand disease and other inherited bleeding disorders in women with diagnosed menorrhagia

Dilley, A. / Drews, C. / Miller, C. / Lally, C. / Austin, H. / Ramaswamy, D. / Lurye, D. / Evatt, B., Obstetrics and Gynecology, Apr 2001

...is a common symptom of **von Willebrand disease**. 8,9 **von Willebrand disease** is a genetic condition...type III) or quality (**type II**) of von Willebrand factor...disorders, most of which were **von Willebrand disease**, in 17 and 37% of women...

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	18.	Investigation of type IIC von Willebrand disease.  H Uno / N Nishida / J Ishizaki / M Suzuki / T Nishikubo / S Miyata Y Takahashi / () / K Tsuda, Int J Hematol, Apr 1994  Type II von Willebrand disease (vWD) is characterized by qualitative abnormality of von Willebrand factor (vWF). It is characterized by the absence of the largest
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		Studies of vWF-botrocetin complex in patients with different subtypes of type ii von willebrand disease - F.I.Pareti,  Thrombosis Research, Jan 1993
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S1	1	("5321127").PN.	USPAT; USOCR	OR	OFF	2006/03/25 11:37
S3	29289	His with val	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 11:39
S4	0	S3 same GP1b\$2	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 11:44
S5	74	glycocalicin	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 11:45
S8	142491	latex	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 14:34
S7	17	S6 and (ELISA RIA)	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 14:34
S12	4	S11 and S8	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 14:38
S13	31	S8 same S9	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 15:44
S17	100	S15 same S16	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 15:45
S16	587401	GP\$3	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 15:45

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S15	834	ristocetin	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 15:45
S6	30	S5 and ristocetin	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 15:45
S20	1	("4935339").PN.	USPAT; USOCR	OR	OFF	2006/03/25 16:31
S21	305019	(agglutinat\$5 aggregat\$6)	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 16:48
S11	76	S9 same ristocetin same Gp\$3	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 16:48
S10	152	S9 same ristocetin and Gp\$3	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 16:48
S9	22530	platelet with (agglutinat\$5 aggregat\$6)	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 16:48
S22	5286	(von with willebrand with factor vWF)	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 16:49
S18	60	S17 and (von with willebrand with disease vWD)	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 16:49
S14	1	("5231033").PN.	USPAT; USOCR	OR	OFF	2006/03/25 16:49
S26	24	S23 and S25	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 16:50

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S25	142491	latex	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 16:50
S23	373	S21 same Gp\$3 same S22	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/25 16:50
S19	13	S17 same (von with willebrand with disease vWD)	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/27 11:22
L5	22532	platelet with (agglutinat\$5 aggregat\$6)	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/27 11:23
L7	1521	latex same ELISA	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/27 11:30
L6	15	L5 same ((light near5 scattering) turbidimetr\$3 ((electric magentic) adj2 field)) with detect\$3	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/27 11:30
L8	771	latex near3 (bead particle)same ELISA	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2006/03/27 11:31

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